CAREONE ANTIBACTERIAL HAND SANITIZER FESTIVAL OF LIGHTS- ethyl alcohol liquid
AMERICAN SALES COMPANY

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

ACTIVE INGREDIENT

ETHYL ALCOHOL 65%

PURPOSE

ANTISEPTIC

USES

TO DECREASE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY. FLAMMABLE. KEEP AWAY FROM SOURCE OF HEAT OR FIRE

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE WITH WATER

STOP USE AND ASK A DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

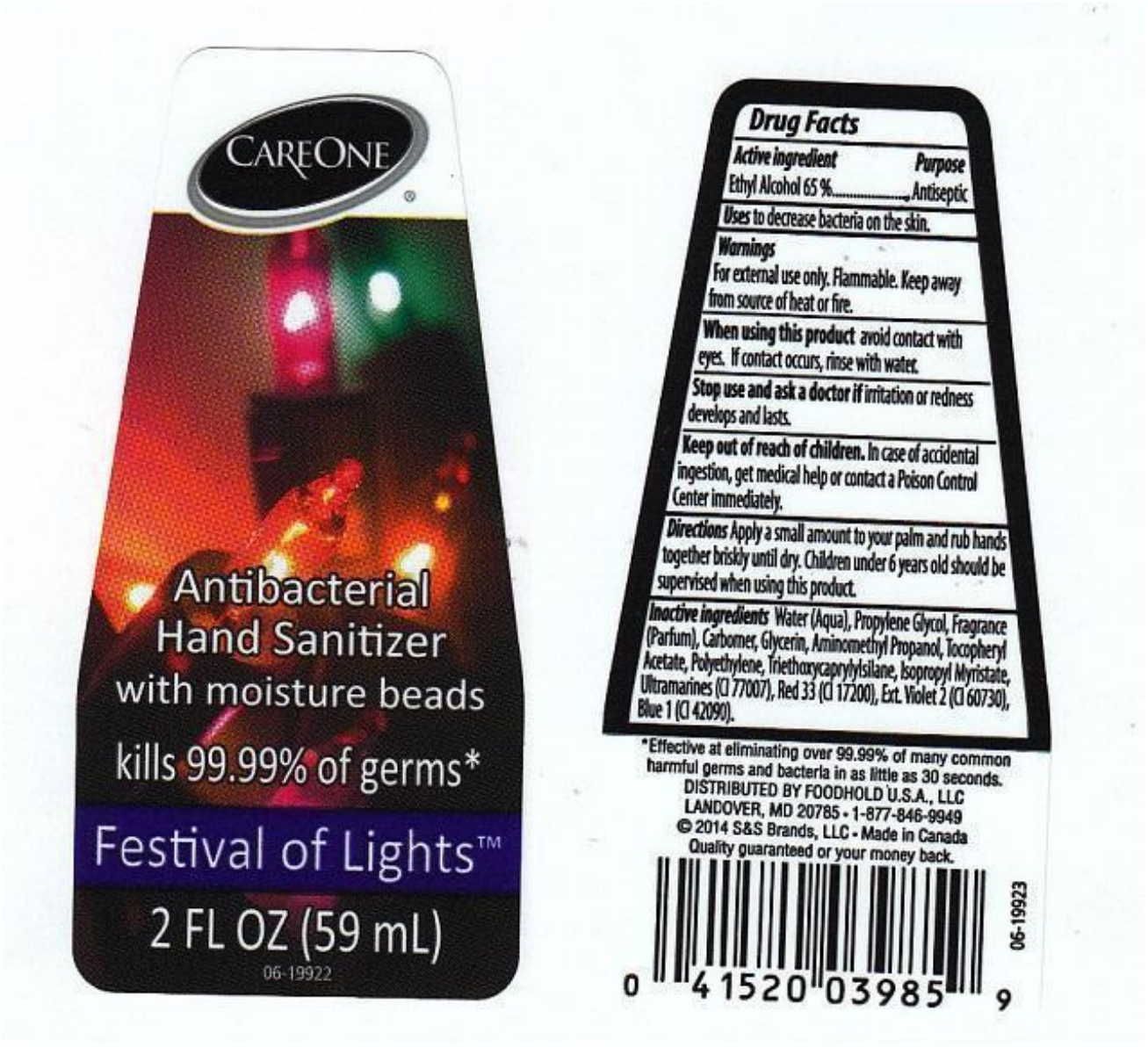
DIRECTIONS

APPLY A SMALL AMOUNT TO YOUR PALM AND RUB HANDS TOGETHER BRISKLY UNTIL DRY. CHILDREN UNDER 6 YEARS OLD SHOULD BE SUPERVISED WHEN USING THIS PRODUCT

INACTIVE INGREDIENT

WATER (AQUA), PROPYLENE GLYCOL, FRAGRANCE (PARFUM), CARBOMER, GLYCERIN, AMINOMETHYL PROPANOL, TOCOPHERYL ACETATE, POLYETHYLENE, TRIETHOXYCAPRYLYLSILANE, ISOPROPYL MYRISTATE, ULTRAMARINES (CI 77007), RED 33 (CI 17200), EXT. VIOLET 2 (CI 60730), BLUE 1 (CI 42090)

LABEL COPY



CAREONE ANTIBACTERIAL HAND SANITIZER FESTIVAL OF LIGHTS

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41520-502
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	650 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER 934 (UNII: Z135WT9208)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ULTRAMARINE BLUE (UNII: I39WR998BI)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41520-502-02	59 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/14/2014	

Labeler - AMERICAN SALES COMPANY (809183973)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(41520-502)

Revised: 8/2014 AMERICAN SALES COMPANY